

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: December 15, 2003, 14:50:12 ; Search time 42 Seconds
(without alignments)
1171.553 Million cell upd

Title: US-09-831-805A-6
 Perfect score: 1635
 Sequence: 1 MALLRRPDLRLCARLPDFEL.....VNYVIRDEEGDFPHKKSSEVT 310

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues
Total number of hits satisfying chosen parameters: 1107863

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000
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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_19Jun03.*

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24:	/SIDSL1/gcgdata/geneseq/geneseq-exp-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Result No.	Score	Query %		Length	DB	ID	Description
		Match	Length				
1	1635	100.0	310	21	AA956294	Human IGFAM-6 immu	
2	1629	99.6	310	21	AAB27276	Human confluency r	
3	1629	99.6	310	21	AAB33457	Human PRO1868 prot	
4	1629	99.6	310	21	AA956735	PRO1868, an A33 an	
5	1629	99.6	310	22	AAAG33323	Human polypeptide,	
6	1629	99.6	310	22	AAAG31905	Human polypeptide,	
7	1629	99.6	310	22	AAU12440	Human PRO1868 poly	
8	1629	99.6	310	22	AAB80272	Human PRO1868 prot	
9	1629	99.6	310	22	AAB80383	Secreted protein e	

10	1629	99.6	310	22	AA80408	Secreted protein e
11	1629	99.6	310	22	AA80409	Secreted protein e
12	1629	99.6	310	23	ABG31401	Human PRO1868 poly
13	1629	99.6	310	23	ABG91361	Novel human secret
14	1629	99.6	310	23	ABG92709	Human secreted pro
15	1629	99.6	310	23	ABG65296	Human albumin fusi
16	1629	99.6	310	23	ABG65297	Human albumin fusi
17	1629	99.6	310	23	ABG65298	Human albumin fusi
18	1629	99.6	310	23	ABG95553	Human angiogenesis
19	1629	99.6	310	23	ABX84947	Human PRO1868 prot
20	1629	99.6	310	24	ABU69682	Novel human secret
21	1629	99.6	310	24	ABU71505	Human PRO polypept
22	1629	99.6	310	24	ABU71951	Human secreted/tr
23	1629	99.6	310	24	ABU66838	Human PRO polypept
24	1629	99.6	310	24	ABU67114	Human secreted/tr
25	1629	99.6	310	24	ABU67405	Human secreted pro
26	1629	99.6	310	24	ABU59919	Novel secreted and
27	1629	99.6	310	24	ABU60813	Human secreted/tr
28	1629	99.6	310	24	ABU64559	Human secreted/tr
29	1629	99.6	310	24	ABG73314	Human PRO1868 poly
30	1629	99.6	310	24	ABP71277	Human functional a
31	1629	99.6	310	24	ABU54407	Human secreted/tr
32	1629	99.6	311	21	AA838333	Human secreted pro
33	1629	99.6	311	21	AA838383	Human secreted pro
34	1629	99.6	311	21	AA838394	Human secreted pro
35	1629	99.6	319	22	AA880431	Gene #13 associate
36	1629	99.1	329	23	ABF41902	Human ovarian anti
37	1613	98.7	310	24	AAO16453	Human functional a
38	1490	91.1	285	21	AA839254	Human secreted pro
39	1475	90.2	321	23	ABG06037	Human NS protein s
40	1401	85.7	310	21	AA827272	Human confluency r
41	1401	85.7	310	21	AA827278	Murine confluency
42	1066	65.2	361	22	ABG22401	Novel human diagno
43	878	53.7	291	22	ABG04645	Novel human diagno
44	878	53.7	404	22	ABG12109	Novel human diagno
45	488	29.8	298	24	AA016452	Human functional a

ALIGNMENTS

RESULT 1	
AA96294	
ID	AA96294 standard; protein; 310 AA.
XX	
XX	AA96294;
XX	
DT	16-AUG-2000 (first entry)
XX	
DE	Human IGFAM-6 immunoglobulin.
XX	
KW	Human; immunoglobulin; IGFAM-6; IGFAM; immune disorder; cancer;
KW	infection; inflammation; haematopoiesis; AIDS; allergy.
XX	
OS	Homo sapiens.

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XX 19-NOV-1998; 99US-0113635.
PR 22-DEC-1998; 98US-0113635.
PR 07-APR-1999; 99US-0128194.
XX (INCY-) INCYTE PHARM INC.
PA Yue H, Tang YT, Corley NC, Guegler KJ, Gorgone GA, Baughn MR;
XX Lu DAM, Lal P, Hillman JL, Yang J;
XX WPI; 2000-387796/33.
DR N-PSDB; AAA27386.
XX Immunoglobulin superfamily proteins, the agonist and antagonist of the
PT protein is useful for preventing and treating disorders associated with
PT altered levels of the protein such as cancer, immune system disorders
PT -
XX Claim 1; Page 82-83; 105pp; English.
PS The present sequence is the human immunoglobulin superfamily protein
XX IGFAM-6. Its gene was isolated from a cDNA library of leg
CC tissue. It is expressed in reproductive, nervous and
CC cardiovascular tissue, where cancer and inflammation are common. The
CC gene, protein, its antibodies, agonists and antagonists are suitable for
CC diagnosing and treating many diseases, including cancer, immune system
CC disorders (such as inflammation, AIDS, allergies, anaemia, Crohn's
CC arteriosclerosis, asthma, atherosclerosis, cholecystitis, Crohn's
CC disease, diabetes mellitus, emphysema, Graves' disease, hepatitis,
CC multiple sclerosis, psoriasis, rheumatoid arthritis, scleroderma,
CC systemic lupus erythematosus and ulcerative colitis), complications of
CC cancer, haemodialysis and extracorporeal circulation, trauma and
CC haematopoietic cancer (such as leukaemia) and infections caused by
CC bacteria, viruses, fungi or parasites.
XX SQ Sequence 310 AA;
Query Match 100.0%; Score 1635; DB 21; Length 310;
Best Local Similarity 100.0%; Pred. No. 2.2e-133;
Matches 310; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVLTQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVLTQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
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QY 241 NIGGIIGVLVAVLALITIGICAYRRGYFINNKQGESYKPKGPDGVNYIRTBEG 300
Db 241 NIGGIIGVLVAVLALITIGICAYRRGYFINNKQGESYKPKGPDGVNYIRTBEG 300
QY 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310
RESULT 2
AAB27276
ID AAB27276 standard; Protein; 310 AA.
XX AAB27276;
XX 23-FEB-2001 (first entry)

```

Human confluency regulated adhesion molecule 1 #2.

Immunoglobulin superfamily; Ig Sf; vascular adhesion molecule; inflammation; cancer; wound; angiogenesis; human; confluency regulated adhesion molecule 1; CRAM-1; JAM-2.

Homo sapiens.

WO200053749-A2.

14-SEP-2000.

13-MAR-2000; 2000WO-EP02219.

11-MAR-1999; 99EP-0200746.

(RMFD-) RMF DICTAGENE SA.

Imhof BA, Aurrand-Lions M;

WPI; 2000-587436/55.

N-PSDB; AAA95306.

Isolated human Confluency Regulated Adhesion Molecule 1 or 2 (CRAM-1 or CRAM-2) polypeptide, useful for treatment of tumors, inflammation reactions and modulating vascular permeability -

Claim 2; Fig 6; 59pp; English.

The present sequence is the human confluency regulated adhesion molecule 1 (CRAM-1, also known as JAM-2). CRAM-1 is one of the vascular adhesion proteins of the immunoglobulin superfamily (Ig Sf). The CRAM-1 protein and coding sequence can be used in the treatment of cancer, inflammation, to modulate cell-cell interactions and angiogenesis, and in the modulation of wound healing.

Query Match 99.6%; Score 1629; DB 21; Length 310; Best Local Similarity 99.7%; Pred. No. 7.2e-133; Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60

Db 1 MALRRPRLRLCARLPDFFLLLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60

QY 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120

Db 61 SDPRIWKKIODEQTTVFFDNKIQDLAGRAEILGKTSLKINWTRRDSALYRCEVVAR 120

QY 121 NDRKEIDEIVLTQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180

Db 121 NDRKEIDEIVLTQVKPVTVCVRKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180

QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQEVEVDL 240

Db 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQEVEVDL 240

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Db 241 NIGGIIGVLVAVLALITIGICAYRRGYFINNKQGESYKPKGPDGVNYIRTBEG 300

QY 301 DFRHKSSFVI 310

Db 301 DFRHKSSFVI 310

RESULT 3

AAB33457

ID AAB33457 standard; Protein; 310 AA.

XX AAB33457;

XX DT 29-JAN-2001 (first entry)

XX DE Human PRO1868 protein UNQ859 SEQ ID NO:193.

XX KW Human; immune related disease; diagnosis; antiinflammatory; cardiant;

XX KW dermatological; antiarthritic; antirheumatic; immunosuppressive;

XX KW haemostatic; antithyroid; antidiabetic; nootropic; neuroprotective;

XX KW antianemic; hepatotropic; virucide; antipsoriatic; antiallergic;

XX KW antiasthmatic; systemic lupus erythematosus; rheumatoid arthritis;

XX KW osteoarthritis; spondyloarthropathy; systemic sclerosis; sarcoidosis;

XX KW idiopathic inflammatory myopathy; Sjogren's syndrome; thyroiditis;

XX KW systemic vasculitis; autoimmune haemolytic anaemia; diabetes mellitus;

XX KW autoimmune thrombocytopenia; immune-mediated renal disease;

XX KW demyelinating disease; hepatobiliary disease; Whipple's disease;

XX KW inflammatory bowel disease; gluten-sensitive enteropathy;

XX KW autoimmune disease; immune-mediated skin disease; allergic disease;

XX KW immunological disease; transplantation associated disease;

XX KW graft rejection; graft-versus-host-disease.

OS Homo sapiens.

XX KW WO200053759-A2.

XX PD 14-SEP-2000.

XX PF 02-MAR-2000; 2000WO-US05841.

XX PR 08-MAR-1999; 99WO-US05028.

XX PR 10-MAR-1999; 99US-0123618.

XX PR 12-MAR-1999; 99US-0123957.

XX PR 23-MAR-1999; 99US-0125775.

XX PR 12-APR-1999; 99US-0128849.

XX PR 20-APR-1999; 99WO-US08615.

XX PR 28-APR-1999; 99US-0134445.

XX PR 04-MAY-1999; 99US-0132371.

XX PR 14-MAY-1999; 99US-0134287.

XX PR 02-JUN-1999; 99WO-US12252.

XX PR 23-JUN-1999; 99US-0141037.

XX PR 20-JUL-1999; 99US-0144758.

XX PR 26-JUL-1999; 99US-0145698.

XX PR 28-JUL-1999; 99US-0146222.

XX PR 01-SEP-1999; 99WO-US20111.

XX PR 08-SEP-1999; 99WO-US20594.

XX PR 13-SEP-1999; 99WO-US20944.

XX PR 15-SEP-1999; 99WO-US21090.

XX PR 15-SEP-1999; 99WO-US21547.

XX PR 05-OCT-1999; 99WO-US23089.

XX PR 29-OCT-1999; 99US-0162506.

XX PR 29-NOV-1999; 99WO-US28214.

XX PR 30-NOV-1999; 99WO-US28313.

XX PR 30-NOV-1999; 99WO-US28409.

XX PR 01-DEC-1999; 99WO-US28301.

XX PR 01-DEC-1999; 99WO-US28634.

XX PR 02-DEC-1999; 99WO-US28551.

XX PR 02-DEC-1999; 99WO-US28564.

XX PR 02-DEC-1999; 99WO-US28565.

XX PR 16-DEC-1999; 99WO-US30095.

XX PR 20-DEC-1999; 99WO-US30999.

XX PR 30-DEC-1999; 99WO-US31274.

XX PR 05-JAN-2000; 2000WO-US00219.

XX PR 06-JAN-2000; 2000WO-US00277.

XX PR 11-FEB-2000; 2000WO-US00376.

XX PR 18-FEB-2000; 2000WO-US03565.

XX PR 18-FEB-2000; 2000WO-US04341.

XX PR 22-FEB-2000; 2000WO-US04342.

XX PR 22-FEB-2000; 2000WO-US04414.

XX PA (GETH) GENENTECH INC.

XX PI Ashkenazi AJ, Baker KP, Goddard A, Gurney AL, Hebert C, Henzel W;

XX PI Kabakoff RC, Lu Y, Pan J, Pennica D, Shelton DL, Smith V;

XX PI Stewart TA, Tumas D, Watanabe CK, Wood WI, Yan M;

XX WI; 2000-572271/53.

XX N-PSDB; AAC58622.

XX PT Sixty four PRO polypeptides, useful in the diagnosis and treatment of

XX PT immune related disorders, e.g. systemic lupus erythematosus, rheumatoid

XX PT arthritis, osteoarthritis, thyroiditis and diabetes mellitus -

XX PS Claim 33; Fig 88; 309pp; English.

XX CC The present invention describes sixty four human PRO proteins which can

XX CC be used in the treatment of immune related diseases. The human PRO

XX CC proteins, anti-PRO antibodies, agonists and antagonists are useful for

XX CC treating and diagnosing immune related disorders. The disorders are

XX CC selected from systemic lupus erythematosus, rheumatoid arthritis,

XX CC osteoarthritis, juvenile chronic arthritis, spondyloarthropathies,

XX CC systemic sclerosis, idiopathic inflammatory myopathies, Sjogren's

XX CC syndrome, systemic vasculitis, sarcoidosis, autoimmune haemolytic

XX CC anaemia, autoimmune thrombocytopenia, thyroiditis, diabetes mellitus,

XX CC immune-mediated renal disease, demyelinating diseases of the central

XX CC and peripheral nervous systems, hepatobiliary diseases, inflammatory

XX CC bowel disease, gluten-sensitive enteropathy and Whipple's disease,

XX CC autoimmune or immune-mediated skin diseases, allergic diseases,

XX CC immunological diseases of the lung, and transplantation associated

XX CC diseases including graft rejection and graft-versus-host-disease.

XX CC AAC58397 to AAC58578 represent PCR primers and hybridisation probes used

XX CC in the isolation of human PRO sequences. AAC58579 to AAC58642 and

XX CC AAB33414 to AAB33477 represent human PRO polynucleotide and protein

XX CC sequences given in the exemplification of the present invention.

XX SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 21; Length 310;

Best Local Similarity 99.7%; Pred. No. 7.2e-133; Indels 0; Gaps 0;

Matches 309; Conservative 0; Mismatches 1;

QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRPVVOEFESVELSCIITDSQT 60

DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRPVVOEFESVELSCIITDSQT 60

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DB 61 SDPRIWKKIODEQTTVFFDNKIQGLAGRAEILGKTSKIWNVTRRDSALYRCEVVAR 120

QY 121 NDRKEIDEIVIELTVQVKPVPVCRKAVPVGKMATLHCQESGHPHPSYWRNDVPL 180

DB 121 NDRKEIDEIVIELTVQVKPVPVCRKAVPVGKMATLHCQESGHPHPSYWRNDVPL 180

QY 181 PTDSRANPRFRNSSSHLNSETGLTVFAVHKDDSGQYVCIASNDAGSARCEQEMEVYDL 240

DB 181 PTDSRANPRFRNSSSHLNSETGLTVFAVHKDDSGQYVCIASNDAGSARCEQEMEVYDL 240

QY 241 NIGGIIGVLVLAVALITLIGICCAVRGYPFINKQDGESYKPKPGDGWNYRTDEEG 300

DB 241 NIGGIIGVLVLAVALITLIGICCAVRGYPFINKQDGESYKPKPGDGWNYRTDEEG 300

QY 301 DFRHKSSRFVI 310

DB 301 DFRHKSSRFVI 310

RESULT 4

AA96735

ID AA96735 standard; Protein; 310 AA.

XX AC AA96735;

XX DT 26-SEP-2000 (first entry)

XX DE PRO1868, an A33 antigen homologue.

XX KW PRO1868; A33 antigen; secreted protein; transmembrane protein;

XX KW anti-inflammatory; cytostatic; recombinant production; gene therapy.

XX OS Homo sapiens.
 XX FH Key
 XX FT Peptide
 XX FT /label= Signal_peptide
 XX FT Modified-site
 XX FT 26..31
 XX FT /note= "N-myristoylation site"
 XX FT Modified-site
 XX FT 69..77
 XX FT /note= "Tyrosine kinase phosphorylation site"
 XX FT Modified-site
 XX FT 104..107
 XX FT /note= "N-glycosylation site"
 XX FT Modified-site
 XX FT 106..109
 XX FT /note= "Casein kinase II phosphorylation site"
 XX FT Modified-site
 XX FT 107..110
 XX FT /note= "cAMP- and cGMP-dependent protein kinase phosphorylation site"
 XX FT Modified-site
 XX FT 192..195
 XX FT /note= "N-glycosylation site"
 XX FT Modified-site
 XX FT 215..220
 XX FT /note= "N-myristoylation site"
 XX FT Modified-site
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 XX FT /label= Transmembrane_domain
 XX FT Modified-site
 XX FT 243..248
 XX FT /note= "N-myristoylation site"
 XX FT Modified-site
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 XX FT /note= "N-myristoylation site"
 XX FT Modified-site
 XX FT 262..267
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 XX FT Modified-site
 XX FT 296..299
 XX FT /note= "Casein kinase II phosphorylation site"
 XX FT
 XX PN WO200036102-A2.
 XX PD 22-JUN-2000.
 XX PF 01-DEC-1999; 99WO-US28634.
 XX PR 16-DEC-1998; 98US-0112851.
 XX PR 16-DEC-1998; 98US-0113145.
 XX PR 22-DEC-1998; 98US-0113511.
 XX PR 12-JAN-1999; 99US-0115558.
 XX PR 12-JAN-1999; 99US-0115565.
 XX PR 12-JAN-1999; 99US-0115733.
 XX PR 09-FEB-1999; 99US-0119341.
 XX PR 10-FEB-1999; 99US-0119537.
 XX PR 12-FEB-1999; 99US-0119965.
 XX PR 02-JUN-1999; 99WO-US12252.
 XX PA (GETH) GENENTECH INC.
 XX PI Botstein D, Desnoyers L, Ferrara N, Fong S, Gao W, Goddard A;
 XX PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
 XX PI Wood WI;
 XX XX
 XX DR WPI; 2000-431586/37.
 XX DR N-PSDB; AAA51265.
 XX PT Isolated nucleic acid molecule encodes a PRO polypeptide which is a
 XX PT transmembrane polypeptide
 XX XX
 XX PS Claim 1; Fig 14; 154pp; English.
 XX CC This is PRO1868, a putative homologue of A33 antigen, a known
 XX CC colorectal cancer-associated marker. The invention concerns novel
 XX CC secreted and transmembrane proteins, designated PRO polypeptides. The
 XX CC cDNA and gene sequences are useful in the recombinant production of PRO
 XX CC polypeptides, as a hybridization probe to screen libraries to isolate
 XX CC cDNAs with sequence identity to PRO polypeptides or to map the gene
 XX CC encoding the PRO polypeptides and analyzing genetic disorders. The
 XX CC cDNA/gene can also be used to produce transgenic animals useful for the

CC development and screening of therapeutically useful reagents. They can
 CC also be used in gene therapy, e.g. to replace a defective gene.
 XX SQ Sequence 310 AA;
 XX Query Match 99.6%; Score 1629; DB 21; Length 310;
 XX Best Local Similarity 99.7%; Pred. No. 7.2e-133;
 XX Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 MALRRPRLRLCARLPDPFFLLLLFRGCLIGAVNLKSSNRTPVVOEFSEVLSCLITDSQT 60
 Db 1 MALRRPRLRLCARLPDPFFLLLLFRGCLIGAVNLKSSNRTPVVOEFSEVLSCLITDSQT 60
 Qy 61 SDPRIWKIQDEQTTVVFNDKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
 Db 61 SDPRIWKIQDEQTTVVFNDKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
 Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVKAIVPGKMATLHCQESGHPHPRHYSWYRNDVPL 180
 Db 121 NDRKEIDEIVIELTVQVKPVPVCRVKAIVPGKMATLHCQESGHPHPRHYSWYRNDVPL 180
 Qy 181 PTDSRANPRFNSHLSNSETGTLVFTAVHKDDSGQYVCIASNDAGSARCEQMEVYDL 240
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 Db 241 NIGGIIGVLVLAVALILTLGICCAVRRGYFINNKDGSYKPKPGDGVNYIRTDEEG 300
 Qy 301 DFRHKSSFVI 310
 Db 301 DFRHKSSFVI 310
 RESULT 5
 AAM93323
 ID AAM93323 standard; Protein; 310 AA.
 XX AC AAM93323;
 XX DT 06-NOV-2001 (first entry)
 XX DE Human polypeptide, SEQ ID NO: 2845.
 XX KW Human; full length cDNA; cDNA synthesis; oligo-capping.
 XX OS Homo sapiens.
 XX PN EPI130094-A2.
 XX PD 05-SEP-2001.
 XX PF 07-JUL-2000; 2000EP-0114089.
 XX PR 08-JUL-1999; 99JP-0194486.
 XX PR 11-JAN-2000; 2000JP-0118774.
 XX PR 02-MAY-2000; 2000JP-0183765.
 XX PA (HELI-) HELIX RES INST.
 XX PI Ota T, Nishikawa T, Isogai T, Hayashi K, Ishii S, Kawai Y;
 XX PI Wakamatsu A, Sugiyama T, Nagai K, Kojima S, Otsuki T, Koga H;
 XX DR WPI; 2001-524255/58.
 XX DR N-PSDB; AAK94243.
 XX PT 830 Primers useful for synthesizing full length cDNA clones and their
 XX PT use in genetic manipulation -
 XX PS Claim 8; SEQ ID NO 2845; 1380pp + sequence listing; English.
 XX CC The invention relates to primers for synthesizing full length cDNA
 XX CC clones. 830 cDNA molecules encoding a human protein have been

CC isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA
CC molecules have been determined. Primers for synthesizing the full length
CC cDNA are useful for clarifying the function of the protein encoded by
CC the cDNA. The full length clones were obtained by construction of full
CC length enriched cDNA libraries that were synthesised by the oligo-capping
CC method. The primers enable the production of the full length cDNA easily
CC without any special methods. The present sequence is a polypeptide
CC encoded by a full length human cDNA of the invention.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in CD-ROM format directly from EPO.
XX
SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Qy 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
Db 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Qy 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Db 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Qy 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGESYKPNPKGPDGVNVRTDEEG 300
Db 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGESYKPNPKGPDGVNVRTDEEG 300
Qy 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310

RESULT 6
AAM93905
ID AAM93905 standard; Protein; 310 AA.

XX AC AAM93905;

XX DT 06-NOV-2001 (first entry)

XX DE Human polypeptide, SEQ ID NO: 4051.

XX KW Human; full length cDNA; cDNA synthesis; oligo-capping.

XX OS Homo sapiens.

XX PN EP1130094-A2.

XX PD 05-SEP-2001.

XX PF 07-JUL-2000; 2000EP-0114089.

XX PR 08-JUL-1999; 99JP-0194486.

XX PR 11-JAN-2000; 2000JP-0119774.

XX PR 02-MAY-2000; 2000JP-0183765.

XX PA (HELI-) HELIX RES INST.

XX PI Ota T, Nishikawa T, Isogai T, Hayashi K, Ishii S, Kawai Y;

XX PI Wakamatsu A, Sugiyama T, Nagai K, Kojima S, Otsuki T, Koga H;

XX WPI; 2001-524255/58.

XX DR N-PSDB; AAK94867.

XX 830 Primers useful for synthesizing full length cDNA clones and their
PT use in genetic manipulation -
XX
XX Claim 8; SEQ ID NO 4051; 1380pp + sequence listing; English.
XX
CC The invention relates to primers for synthesising full length cDNA
CC clones. 830 cDNA molecules encoding a human protein have been
CC isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA
CC molecules have been determined. Primers for synthesising the full length
CC cDNA are useful for clarifying the function of the protein encoded by
CC the cDNA. The full length clones were obtained by construction of full
CC length enriched cDNA libraries that were synthesised by the oligo-capping
CC method. The primers enable the production of the full length cDNA easily
CC without any special methods. The present sequence is a polypeptide
CC encoded by a full length human cDNA of the invention.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in CD-ROM format directly from EPO.
XX
SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Qy 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
Db 61 SDPRIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCOESGHPHYSWYRNDVPL 180
Qy 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Db 181 PTDSRANPRFRNSSHLSNSETGLVFTAVHKDDSGQYICIASNDAGSARCEQEMEYVDL 240
Qy 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGESYKPNPKGPDGVNVRTDEEG 300
Db 241 NIGGIIGVVLVAVLALITIGICCAVRRGYFINNKQDGESYKPNPKGPDGVNVRTDEEG 300
Qy 301 DFRHKSSFVI 310
Db 301 DFRHKSSFVI 310

RESULT 7

AAM12440

ID AAM12440 standard; Protein; 310 AA.

XX AC AAM12440;

XX DT 24-OCT-2001 (first entry)

XX DE Human PRO1868 polypeptide sequence.

XX KW Human secretory and transmembrane; PRO; mammalian; cancer; lung;

XX KW breast; prostate; cervical; tumour necrosis factor-alpha; TNF-alpha;

XX KW cartilage; ear; proliferation; glucose; free fatty acid; skeletal muscle;

XX KW adipocyte; A-peptide; factor VIIa; gene therapy.

XX OS Homo sapiens.

XX PN WO200140466-A2.

XX PD 07-JUN-2001.

XX PF 01-DEC-2000; 2000WO-US32678.

XX XX

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PR 01-DEC-1999; 99WO-US28301.
PR 01-DEC-1999; 99WO-US28634.
PR 02-DEC-1999; 99WO-US28551.
PR 02-DEC-1999; 99WO-US28564.
PR 02-DEC-1999; 99WO-US28565.
PR 09-DEC-1999; 99US-0170262.
PR 16-DEC-1999; 99WO-US30095.
PR 20-DEC-1999; 99WO-US30911.
PR 20-DEC-1999; 99WO-US30999.
PR 30-DEC-1999; 99WO-US31243.
PR 06-JAN-2000; 2000WO-US00277.
PR 06-JAN-2000; 2000WO-US00376.
PR 11-FEB-2000; 2000WO-US03565.
PR 18-FEB-2000; 2000WO-US04341.
PR 18-FEB-2000; 2000WO-US04342.
PR 22-FEB-2000; 2000WO-US04414.
PR 24-FEB-2000; 2000WO-US04914.
PR 24-FEB-2000; 2000WO-US05004.
PR 01-MAR-2000; 2000WO-US05601.
PR 20-MAR-2000; 2000WO-US07377.
PR 21-MAR-2000; 2000WO-US07532.
PR 30-MAR-2000; 2000WO-US08439.
PR 17-MAY-2000; 2000WO-US13705.
PR 22-MAY-2000; 2000WO-US14042.
PR 30-MAY-2000; 2000WO-US14941.
PR 02-JUN-2000; 2000WO-US15264.
PR 10-NOV-2000; 2000WO-US30873.
XX
XX
PA (GETH ) GENENTECH INC.
XX
XX Baker KP, Beresini M, DeForge L, Desnoyers L, Filvaroff E, Gao W;
PI Gerritsen ME, Goddard A, Godowski PJ, Gurney AL, Sherwood S;
PI Smith V, Stewart TA, Tumas D, Watanabe CK, Wood WI, Zhang Z;
XX
XX WPI; 2001-408281/43.
XX N-PSDB; AAS21512.
XX
PT Isolated, secretory and transmembrane PRO polypeptide used to detect
PT other PRO polypeptides, link bioactive molecules to cells expressing
PT PRO polypeptides, and detect the presence of mammalian tumours e.g.
PT lung, breast, prostate, cervical
XX
XX Claim 12; Fig 538; 813pp; English.
XX
XX AAU12172-AAU12446 represent novel human secretory and transmembrane
CC PRO polypeptides. The PRO polypeptides are useful to detect other
CC PRO polypeptides, to link bioactive molecules to cells expressing
CC PRO polypeptides, to modulate biological activities of cells expressing
CC PRO polypeptides, and to detect the presence of mammalian lung, colon,
CC breast, prostate, rectal, cervical or liver tumours by comparing PRO
CC polypeptide expression in a cell sample to that in a control sample.
CC Some of the 275 sequences are also useful to stimulate the release of
CC tumour necrosis factor-alpha (TNF-alpha) from human blood, the
CC proliferation or differentiation of chondrocytes, the proliferation or
CC gene expression in pericyte cells, the release of proteoglycans from
CC cartilage, the proliferation of inner ear utricular supporting cells or
CC of T-lymphocytes, the release of a cytokine from peripheral blood
CC monocytes (PBMCs), or the proliferation of endothelial cells. Some of
CC the PRO polypeptides may modulate glucose or free fatty acid uptake by
CC skeletal muscle cells or by adipocytes; or inhibit binding of A-peptide
CC to factor VIIA. The PRO polypeptides can be used in assays to identify
CC molecules involved in binding interactions. The polynucleotides encoding
CC PRO polypeptides can be used to generate probes, antisense RNA/DNA,
CC transgenic or knock out animals and can be used in gene therapy.
XX
XX Sequence 310 AA;
SQ
Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
XX
XX 1 MALRRPRLRLCARLPDFFLLFRGLIGAVNLKSSNRTPVVQEFESVLSLCIITDSQT 60
|||||

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Db 1 MALRRPRLRLCARLPDFFLLFRGLIGAVNLKSSNRTPVVQEFESVLSLCIITDSQT 60
Qy 61 SDPIEWKKIQDQTTTVVFDNKKIQDLAGRAEILGKTSLKINWVTRRDSALYRCEVVAR 120
|||||
Db 61 SDPIEWKKIQDQTTTVVFDNKKIQDLAGRAEILGKTSLKINWVTRRDSALYRCEVVAR 120
Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
|||||
Db 121 NDRKEIDEIVIELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Qy 181 PTDSRANPRNRSSSHLNSGTGLVFTAVHKDSSGOYVCIASNDAGSARCEOEEMEYVDL 240
|||||
Db 181 PTDSRANPRNRSSSHLNSGTGLVFTAVHKDSSGOYVCIASNDAGSARCEOEEMEYVDL 240
Qy 241 NIGGIIGGLVLAVALALITLIGICCAVRRGYFINNKQDGESYKPKGPDGVNVRTDEEG 300
|||||
Db 241 NIGGIIGGLVLAVALALITLIGICCAVRRGYFINNKQDGESYKPKGPDGVNVRTDEEG 300
Qy 301 DFRHKSSFVI 310
|||||
Db 301 DFRHKSSFVI 310
XX
XX RESULT 8
XX AAB80272
XX ID AAB80272 standard; Protein; 310 AA.
XX AC AAB80272;
XX
XX 24-APR-2001 (first entry)
XX DT Human PRO1868 protein.
XX
XX Human; PRO; dermatological; antipsoriatic; cytostatic; antiinflammatory;
XX antiparkinsonian nootropic; neuroprotective; vulnerary; cardiant;
XX antiangiogenic; vasotropic; antiasthmatic; antirheumatic; cancer;
XX antiarthritic; antiinfertility; antidiabetic; antiviral; diabetes;
XX ophthalmological; gene therapy; skin disease; gastrointestinal disorder;
XX ischaemia; inflammation.
XX
XX Homo sapiens.
XX
XX WO200104311-A1.
XX
XX 18-JAN-2001.
XX
XX 22-FEB-2000; 2000WO-US04414.
XX
XX 07-JUL-1999; 99US-0143048.
XX 26-JUL-1999; 99US-0145698.
XX 28-JUL-1999; 99US-0146222.
XX 08-SEP-1999; 99WO-US20594.
XX 13-SEP-1999; 99WO-US20944.
XX 15-SEP-1999; 99WO-US21090.
XX 15-SEP-1999; 99WO-US21547.
XX 05-OCT-1999; 99WO-US23089.
XX 29-NOV-1999; 99WO-US28214.
XX 30-NOV-1999; 99WO-US28313.
XX 16-DEC-1999; 99WO-US30095.
XX 20-DEC-1999; 99WO-US30911.
XX 20-DEC-1999; 99WO-US30999.
XX 05-JAN-2000; 99WO-US00219.
XX
XX (GETH ) GENENTECH INC.
XX
XX Ashkenazi AJ, Botstein D, Desnoyers L, Eaton DL, Ferrara N;
XX Filvaroff E, Fong S, Gao W, Gerber H, Gerritsen ME, Goddard A;
XX Godowski PJ, Grimaldi CJ, Gurney AL, Hillan KJ, Kljavin IJ;
XX Mather JP, Pan J, Paoni NF, Roy MA, Stewart TA, Tumas D;
XX Williams PM, Wood WI;
XX
XX WPI; 2001-081051/09.
XX N-PSDB; AAF72433.
XX

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XX Sixty one nucleic acids encoding PRO polypeptides which are useful in
PT the treatment of skin diseases (e.g. psoriasis), cancers (e.g. lung
PT squamous cell carcinoma) and neurodegenerative diseases (e.g.
PT Alzheimer's disease) -
XX
XX Claim 1; Fig 124; 393pp; English.
XX
CC The present sequence is one of sixty one novel secreted and
CC transmembrane PRO polypeptides. The PRO polypeptides are
CC useful for treating skin diseases (e.g. psoriasis), cancers (e.g. lung
CC squamous cell carcinoma), gastrointestinal disorders (e.g.
CC enterocolitis), neurodegenerative diseases (e.g. Alzheimer's disease,
CC Parkinson's disease), wound repair, cardiovascular disorders (e.g.
CC endometrial bleeding angiogenesis, ischaemias such as coronary
CC ischaemia, atherosclerosis), inflammatory disorders (e.g. asthma,
CC rheumatoid arthritis, multiple sclerosis), infertility, AIDS and
CC diabetes and retinal disorders such as retinitis pigmentosum.
CC The PRO nucleic acids have applications in molecular biology, including
CC use as hybridization probes, and in chromosome and gene mapping.
XX
SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60

QY 61 SDPIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTRRDSALYRCEVVAR 120
DB 61 SDPIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTRRDSALYRCEVVAR 120

QY 121 NDRKEIDEIVLTVOVKPVPVCVRKAVPVGKMATLHCOSEGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVLTVOVKPVPVCVRKAVPVGKMATLHCOSEGHPRPHYSWYRNDVPL 180

QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQMEVYDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQMEVYDL 240

QY 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300

QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310

RESULT 9
AAB80383
ID AAB80383 standard; protein; 310 AA.
XX
XX AAB80383;
XX
DT 24-APR-2001 (first entry)
XX
DE Secreted protein encoded by gene #13.
XX
KW Secreted protein; human; autoimmune; hyperproliferation;
KW cardiovascular; cerebrovascular; infection; food.
XX
OS Homo sapiens.
XX
PN WO200107459-A1.
XX
PD 01-FEB-2001.
XX
PF 20-JUL-2000; 2000WO-US19735.
XX

PR 23-JUL-1999; 99US-0145220.
XX (HUMA-) HUMAN GENOME SCI INC.
PA
XX
PI Rosen CA, Ruben SM, Ebner R, Duan RD, Ni J, Soppet DR, Moore PA;
PI Shi Y, Lafleur DW, Olsen HS, Birse CE, Komatsoulis GA;
XX
XX WPI; 2001-123261/13.
XX
XX New isolated nucleic acid encoding 29 secreted proteins, for
PT diagnosing, preventing and treating e.g. autoimmune,
PT hyperproliferative, cardiovascular, and ocular diseases or disorders
PT and microorganism infections -
PS
XX Claim 11; Page 538-539; 601pp; English.
XX
CC The present invention relates to 29 human secreted proteins. The
CC invention is used to prevent autoimmune diseases e.g. rheumatoid
CC arthritis, hyperproliferative disorders e.g. neoplasms of the
CC breast or liver, cardiovascular disorders e.g. cardiac arrest,
CC cerebrovascular disorders e.g. cerebral ischemia, angiogenesis,
CC nervous system disorders e.g. Alzheimer's disease, infections
CC caused by bacteria, viruses and fungi and ocular disorders e.g.
CC corneal infection. Also used in food preparations.
XX
SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 22; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLFRGCLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60

QY 61 SDPIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTRRDSALYRCEVVAR 120
DB 61 SDPIEWKKIQDEQTYVFFDNKIQDLAGRAEILGKTSLKIMNVTRRDSALYRCEVVAR 120

QY 121 NDRKEIDEIVLTVOVKPVPVCVRKAVPVGKMATLHCOSEGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVLTVOVKPVPVCVRKAVPVGKMATLHCOSEGHPRPHYSWYRNDVPL 180

QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQMEVYDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGYYCIASNDAGSARCEQMEVYDL 240

QY 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGVLVLAVALITLIGICCAVRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300

QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310

RESULT 10
AAB80408
ID AAB80408 standard; protein; 310 AA.
XX
XX AAB80408;
XX
DT 24-APR-2001 (first entry)
XX
DE Secreted protein encoded by gene #38.
XX
KW Secreted protein; human; autoimmune; hyperproliferation;
KW cardiovascular; cerebrovascular; infection; food.
XX
OS Homo sapiens.
XX
PN WO200107459-A1.
XX

```
PD 01-FEB-2001.
XX
XX
XX 20-JUL-2000; 2000WO-US19735.
XX
XX 23-JUL-1999; 99US-0145220.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM, Ebner R, Duan RD, Ni J, Soppet DR, Moore PA;
XX Shi Y, Lafleur DW, Olsen HS, Birse CE, Komatsoulis GA;
XX WPI; 2001-123261/13.
XX
XX New isolated nucleic acid encoding 29 secreted proteins, for
XX diagnosing, preventing and treating e.g. autoimmune,
XX hyperproliferative, cardiovascular, and ocular diseases or disorders
XX and microorganism infections -
XX
XX Claim 11; Page 557-558; 601pp; English.
XX
XX The present invention relates to 29 human secreted proteins. The
XX invention is used to prevent autoimmune diseases e.g. rheumatoid
XX arthritis, hyperproliferative disorders e.g. neoplasms of the
XX breast or liver, cardiovascular disorders e.g. cardiac arrest,
XX cerebrovascular disorders e.g. cerebral ischemia, angiogenesis,
XX nervous system disorders e.g. Alzheimer's disease, infections
XX caused by bacteria, viruses and fungi and ocular disorders e.g.
XX corneal infection. Also used in food preparations.
XX
XX Sequence 310 AA;
XX
XX Query Match 99.6%; Score 1629; DB 22; Length 310;
XX Best Local Similarity 99.7%; Pred. No. 7.2e-133;
XX Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
QY 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
XX
QY 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
XX
QY 121 NDRKEIDEIVIELTVQVKPVTVCVRPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVTVCVRPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
XX
QY 181 PTDSRANPRFRNSSSHLNSETGTLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
Db 181 PTDSRANPRFRNSSSHLNSETGTLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
XX
QY 241 NIGGIIGGLVWLAVLALITLIGICAYRRGYFINNKQDGESYKPNKPGDGVNIRTDEEG 300
Db 241 NIGGIIGGLVWLAVLALITLIGICAYRRGYFINNKQDGESYKPNKPGDGVNIRTDEEG 300
XX
QY 301 DFRHKSSFFVI 310
Db 301 DFRHKSSFFVI 310
XX
RESULT 11
AAB80409
ID AAB80409 standard; protein; 310 AA.
XX
XX AAB80409;
AC
XX
XX 24-APR-2001 (first entry)
XX
XX Secreted protein encoded by gene #39.
XX
XX Secreted protein; human; autoimmune; hyperproliferation;
KW cardiovascular; cerebrovascular; infection; food.
XX
XX
```

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OS Homo sapiens.
XX
XX WO200107459-A1.
XX
XX 01-FEB-2001.
XX
XX 20-JUL-2000; 2000WO-US19735.
XX
XX 23-JUL-1999; 99US-0145220.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM, Ebner R, Duan RD, Ni J, Soppet DR, Moore PA;
XX Shi Y, Lafleur DW, Olsen HS, Birse CE, Komatsoulis GA;
XX WPI; 2001-123261/13.
XX
XX New isolated nucleic acid encoding 29 secreted proteins, for
XX diagnosing, preventing and treating e.g. autoimmune,
XX hyperproliferative, cardiovascular, and ocular diseases or disorders
XX and microorganism infections -
XX
XX Claim 11; Page 559-560; 601pp; English.
XX
XX The present invention relates to 29 human secreted proteins. The
XX invention is used to prevent autoimmune diseases e.g. rheumatoid
XX arthritis, hyperproliferative disorders e.g. neoplasms of the
XX breast or liver, cardiovascular disorders e.g. cardiac arrest,
XX cerebrovascular disorders e.g. cerebral ischemia, angiogenesis,
XX nervous system disorders e.g. Alzheimer's disease, infections
XX caused by bacteria, viruses and fungi and ocular disorders e.g.
XX corneal infection. Also used in food preparations.
XX
XX Sequence 310 AA;
XX
XX Query Match 99.6%; Score 1629; DB 22; Length 310;
XX Best Local Similarity 99.7%; Pred. No. 7.2e-133;
XX Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
QY 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
Db 1 MALRRPRLRLCARLPDPFLLFRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
XX
QY 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
Db 61 SDPRIWKKIODEQTYVFFDNKIQDLAGRAEILGKTSKINWTRRDSALYRCEVVAR 120
XX
QY 121 NDRKEIDEIVIELTVQVKPVTVCVRPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
Db 121 NDRKEIDEIVIELTVQVKPVTVCVRPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
XX
QY 181 PTDSRANPRFRNSSSHLNSETGTLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
Db 181 PTDSRANPRFRNSSSHLNSETGTLVFTAVHKDDSGQYICIASNDAGSARCEQEVEVDL 240
XX
QY 241 NIGGIIGGLVWLAVLALITLIGICAYRRGYFINNKQDGESYKPNKPGDGVNIRTDEEG 300
Db 241 NIGGIIGGLVWLAVLALITLIGICAYRRGYFINNKQDGESYKPNKPGDGVNIRTDEEG 300
XX
QY 301 DFRHKSSFFVI 310
Db 301 DFRHKSSFFVI 310
XX
RESULT 12
AAB31401
ID AAB31401 standard; protein; 310 AA.
XX
XX AAB31401;
AC
XX
XX 29-NOV-2002 (first entry)
XX
XX Human PRO1868 polypeptide.
XX
XX
```


XX Human; secreted and transmembrane polypeptide; PRO polypeptide;
 KW T-lymphocyte proliferation; inflammatory disease; rheumatoid arthritis;
 KW inflammatory bowel disease; Sjogren's syndrome; thyroiditis;
 KW autoimmune haemolytic anaemia; diabetes mellitus; multiple sclerosis;
 KW hepatitis; contact dermatitis; allergic disease; psoriasis; virucide;
 KW immune related disease; kidney disease; antinflammatory; antithyroid;
 KW antirheumatic; antiarthritic; immunosuppressive; antianaemic;
 KW antidiabetic; neuroprotective; hepatotropic; antiinflammatory;
 KW dermatological; antiallergic; antipsoriatic; PRO1868.
 XX
 OS Homo sapiens.
 XX

XX Key Location/Qualifiers
 FT Peptide 1..30
 FT /label= Signal_peptide
 FT Modified-site 26..31
 FT /note= "N-myristoylation site"
 FT Protein 31..310
 FT /label= Mature_PRO1868
 FT Modified-site 69..77
 FT /note= "Tyrosine kinase phosphorylation site"
 FT Modified-site 104..107
 FT /note= "N-glycosylation site"
 FT Modified-site 106..109
 FT /note= "Casein kinase II phosphorylation site"
 FT Modified-site 107..110
 FT /note= "cAMP- and cGMP-dependent protein kinase phosphorylation site"
 FT Modified-site 192..195
 FT /note= "N-glycosylation site"
 FT Modified-site 215..220
 FT /note= "N-myristoylation site"
 FT Modified-site 226..231
 FT /note= "N-myristoylation site"
 FT Domain 243..263
 FT /label= Transmembrane_domain
 FT Modified-site 243..248
 FT /note= "N-myristoylation site"
 FT Modified-site 244..249
 FT /note= "N-myristoylation site"
 FT Modified-site 262..267
 FT /note= "N-myristoylation site"
 FT Modified-site 296..299
 FT /note= "Casein kinase II phosphorylation site"
 XX
 US2002098507-A1.
 XX
 PD 25-JUL-2002.
 XX
 XX 27-DEC-2001; 2001US-0033326.
 XX
 PR 02-JUN-1999; 99WO-US12252.
 PR 01-DEC-1999; 99WO-US28634.
 PR 02-DEC-1999; 99WO-US28551.
 PR 11-FEB-2000; 2000WO-US03565.
 PR 02-FEB-2000; 2000WO-US04414.
 PR 02-MAR-2000; 2000WO-US05841.
 PR 30-MAR-2000; 2000WO-US08439.
 PR 30-MAY-2000; 2000WO-US14941.
 PR 02-JUN-2000; 2000WO-US15264.
 PR 01-DEC-2000; 2000WO-US32678.
 PR 16-DEC-1998; 98US-113145P.
 PR 22-DEC-1998; 98US-113511P.
 PR 12-JAN-1999; 99US-115558P.
 PR 12-JAN-1999; 99US-115565P.
 PR 09-FEB-1999; 99US-115733P.
 PR 10-FEB-1999; 99US-119341P.
 PR 10-FEB-1999; 99US-119537P.
 PR 12-FEB-1999; 99US-119965P.
 PR 29-OCT-1999; 99US-162506P.
 XX
 XX (GETH) GENENTECH INC. PA

XX Botstein D, Deenoyers L, Ferrara N, Fong S, Gao W, Goddard A;
 PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
 PI Wood WI;
 XX
 DR WPI; 2002-673823/72.
 DR N-PSDB; ABS53477.
 XX
 PT Novel PRO polypeptides and nucleic acids encoding the polypeptides,
 PT useful for preparing a medicament for the treatment of inflammatory and
 PT immune related disorders -
 XX
 PS Claim 12; Fig 14; 125pp; English.
 XX

CC The present invention relates to the isolation of novel human
 CC secreted and transmembrane polypeptides, designated PRO polypeptides,
 CC and the polynucleotide sequences encoding them. The PRO polypeptides
 CC of the invention include PRO1800, PRO539, PRO982, PRO1434, PRO1863,
 CC PRO1917, PRO1868, PRO3434 and PRO1927. The PRO polypeptides can
 CC inhibit the stimulation of T-lymphocyte proliferation. The PRO
 CC polypeptides are useful for the diagnosis and treatment of inflammatory
 CC diseases (e.g. inflammatory bowel disease, rheumatoid arthritis,
 CC Sjogren's syndrome, autoimmune haemolytic anaemia, thyroiditis, diabetes
 CC mellitus, multiple sclerosis, hepatitis, contact dermatitis, allergic
 CC diseases and psoriasis), immune related diseases, and kidney diseases
 CC in humans. The present sequence represents human PRO1868 polypeptide.
 XX

SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 23; Length 310;
 Best Local Similarity 99.7%; Pred. No. 7.2e-133;
 Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 MALRRPRLRLCARLPDPFELLILFRGCLIGAVNLKSSNRTPVQVEFSEVLSCLITDSQT 60
 Db 1 MALRRPRLRLCARLPDPFELLILFRGCLIGAVNLKSSNRTPVQVEFSEVLSCLITDSQT 60

Qy 61 SDPRIEWKKIQDEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120
 Db 61 SDPRIEWKKIQDEQTYVFFDNKIQGLAGRAEILGKTSLKIMNVTTRDSALYRCEVVAR 120

Qy 121 NDRKEIDEIVIELTVQVKPVPVCRVKAPVGVKQATLHCQSEGHPRPHYSWYRNDVPL 180
 Db 121 NDRKEIDEIVIELTVQVKPVPVCRVKAPVGVKQATLHCQSEGHPRPHYSWYRNDVPL 180

Qy 181 PTDSRANPRFRNSSHLNSETGTLVFTAVHKDDSGQYVCYICASNDAGSARCEQMEVYDL 240
 Db 181 PTDSRANPRFRNSSHLNSETGTLVFTAVHKDDSGQYVCYICASNDAGSARCEQMEVYDL 240

Qy 241 NIGGIIGVLVLAVALILITLIGICAYRRGYFINNKQDGSYKPNKPGDGVNYIRTDDEG 300
 Db 241 NIGGIIGVLVLAVALILITLIGICAYRRGYFINNKQDGSYKPNKPGDGVNYIRTDDEG 300

Qy 301 DFRHKSSFVI 310
 Db 301 DFRHKSSFVI 310

RESULT 13
 ABG91361

ID ABG91361 standard; Protein; 310 AA.

XX AC ABG91361;

XX DT 29-NOV-2002 (first entry)

XX DE Novel human secreted protein #7.

XX KW Human; secreted protein; transmembrane protein; gene mapping;
 XX transgenic; immunogenic.

XX OS Homo sapiens.

XX

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PN US2002098505-A1.
XX 25-JUL-2002.
PD 28-DEC-2001; 2001US-0033246.
XX 02-JUN-1999; 99WO-US12252.
PR 01-DEC-1999; 99WO-US28634.
PR 02-DEC-1999; 99WO-US28551.
PR 11-FEB-2000; 2000WO-US03565.
PR 22-FEB-2000; 2000WO-US04414.
PR 02-MAR-2000; 2000WO-US05841.
PR 30-MAR-2000; 2000WO-US08439.
PR 02-JUN-2000; 2000WO-US14941.
PR 01-DEC-2000; 2000WO-US2678.
PR 16-DEC-1998; 98US-113145P.
PR 22-DEC-1998; 98US-113511P.
PR 12-JAN-1999; 99US-115558P.
PR 12-JAN-1999; 99US-115565P.
PR 12-JAN-1999; 99US-115733P.
PR 09-FEB-1999; 99US-119341P.
PR 10-FEB-1999; 99US-119537P.
PR 12-FEB-1999; 99US-119965P.
PR 29-OCT-1999; 99US-162506P.
XX (GETH ) GENENTECH INC.
PA Botstein D, Desnoyers L, Ferrara N, Fong S, Gao W, Goddard A;
PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
PI Wood WI;
XX WPI; 2002-665999/71.
DR N-PSDB; ABS67460.
XX
XX New human secreted and transmembrane (PRO) polypeptides, useful for
PT treating conditions requiring PRO polypeptides, for screening PRO
PT antagonists and agonists useful as drug candidates -
XX
XX Claim 12; Figure 14; 125pp; English.
XX
XX The invention relates to new human secreted and transmembrane proteins
CC (PRO) and nucleic acids of the invention. The polypeptides can be
CC administered therapeutically, especially by expressing encoding
CC polynucleotides, e.g. in therapeutic compositions. They can be used to
CC screen for PRO polypeptide antagonists and agonists useful to identify
CC drug candidates. They can also be used to produce antibodies, useful to
CC detect PRO polypeptides (e.g. diagnostically), purify PRO polypeptides or
CC therapeutically (e.g. as antagonists or to target and/or deliver
CC cytotoxic agents). The polynucleotides are useful therapeutically e.g. to
CC produce antisense sequences to inhibit polypeptide production. They can
CC be used to produce probes and primers useful to detect or isolate
CC sequences encoding PRO polypeptides or similar sequences e.g. variants or
CC sequences from other species. They are also useful for gene mapping and
CC to generate transgenic animals. ABG91355-ABG91363 represent human PRO
CC amino acid sequences of the invention.
XX
XX Sequence 310 AA;
SQ
Query Match 99.6%; Score 1629; DB 23; Length 310;
Best Local Similarity 99.7%; Pred. No. 7, 2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 MALRRPRLRLCARLPDFLLFRGLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFLLFRGLIGAVNLKSSNRTVPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKIODEQTYTFDNKIOGLAGRAEILGKTSLKINWTRDSALYRCVVAR 120
DB 61 SDPRIWKIODEQTYTFDNKIOGLAGRAEILGKTSLKINWTRDSALYRCVVAR 120
QY 121 NDRKEIDEIVIELTVQKVPVPCVRPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVIELTVQKVPVPCVRPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDSDGQYICIASNDAGSARCEQMEVYDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDSDGQYICIASNDAGSARCEQMEVYDL 240
QY 241 NIGGIIGGLVWLAVLALITLIGICCAVRRGYFINNKODGESYKKNPKGPDGVNIRTDEEG 300
DB 241 NIGGIIGGLVWLAVLALITLIGICCAVRRGYFINNKODGESYKKNPKGPDGVNIRTDEEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310
XX
XX RESULT 14
XX ABG92709
XX ID ABG92709 standard; Protein; 310 AA.
XX AC ABG92709;
XX DT 18-NOV-2002 (first entry)
XX DE Human secreted protein PRO1868.
XX KW Human; secreted and transmembrane protein; PRO1800; PRO539;
XX KW PRO982; PRO1434; PRO1863; PRO1917; PRO1868; PRO3434; PRO1927;
XX KW inflammatory disorder; immune related disease; rheumatoid arthritis;
XX KW systemic lupus erythematosus; systemic sclerosis; thyroiditis;
XX KW autoimmune haemolytic anaemia; diabetes mellitus; infectious hepatitis;
XX KW psoriasis; allergic disease of the lung; graft-versus host disease;
XX KW tumour; gene therapy.
XX OS Homo sapiens.
XX PN US2002098506-A1.
XX PD 25-JUL-2002.
XX PF 27-DEC-2001; 2001US-0033301.
XX 04-AUG-1998; 98US-095325P.
XX 16-DEC-1998; 98US-112851P.
XX 16-DEC-1998; 98US-113145P.
XX 22-DEC-1998; 98US-113511P.
XX 12-JAN-1999; 99US-115558P.
XX 12-JAN-1999; 99US-115565P.
XX 09-FEB-1999; 99US-119341P.
XX 10-FEB-1999; 99US-119537P.
XX 12-FEB-1999; 99US-119965P.
XX 29-OCT-1999; 99US-162506P.
XX 02-JUN-1999; 99WO-US12252.
XX 01-DEC-1999; 99WO-US28634.
XX 02-DEC-1999; 99WO-US28551.
XX 11-FEB-2000; 2000WO-US03565.
XX 22-FEB-2000; 2000WO-US04414.
XX 02-MAR-2000; 2000WO-US05841.
XX 30-MAR-2000; 2000WO-US08439.
XX 02-MAY-2000; 2000WO-US14941.
XX 02-JUN-2000; 2000WO-US15264.
XX 01-DEC-2000; 2000WO-US2678.
XX (GETH ) GENENTECH INC.
PA Botstein D, Desnoyers L, Ferrara N, Fong S, Gao W, Goddard A;
PI Gurney AL, Pan J, Roy MA, Stewart TA, Tumas D, Watanabe CK;
PI Wood WI;
XX WPI; 2002-690475/74.
DR N-PSDB; ABS68392.
XX
XX Novel secreted and transmembrane polypeptides and polynucleotides
```

PT useful for diagnosis and treatment of inflammatory disorders and
FI immune-related diseases, and identifying modulators -
XX
PS Claim 12; Fig 14; 125pp; English.

XX The invention relates to an isolated polypeptide having at least 80%
CC amino acid sequence identity to secreted and transmembrane polypeptides
CC PRO1800, PRO539, PRO1434, PRO1863, PRO1868, PRO3434 or
CC PRO1927 and their encoding nucleic acids. Also included are vectors, host
CC cells and antibodies against PRO polypeptides. PRO proteins are useful
CC for identifying modulators of the polypeptide. PRO1868 useful for the
CC diagnosis and treatment of inflammatory and immune related diseases
CC including systemic lupus erythematosus, rheumatoid arthritis, systemic
CC sclerosis, autoimmune haemolytic anaemia, thyroiditis, diabetes mellitus,
CC infectious hepatitis, psoriasis, allergic diseases of the lung and
CC graft-versus host disease and tumours. PRO nucleic acids are useful for
CC constructing hybridisation probes for mapping the gene that encodes that
CC PRO and for the genetic analysis of individuals with genetic disorders,
CC and for generating transgenic animals which are useful in the development
CC and screening of therapeutically useful reagents. PRO nucleic acids are
CC also useful for gene therapy, chromosome identification, and tissue
CC typing. PRO proteins are useful as molecular weight markers for protein
CC electrophoresis purposes. The anti-PRO antibodies are useful in
CC diagnostic assays for PRO, e.g. detecting its expression in specific
CC cells, tissues or serum and for affinity purification of PRO.
XX The present sequence represents a PRO protein.

SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 23; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MALRRPRLRLCARLPDFFLLLRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLLRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKINWVTRRDSALYRCEVVAR 120
DB 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKINWVTRRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYCIASNDAGSARCEQMEVYDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYCIASNDAGSARCEQMEVYDL 240
QY 241 NIGGIIGGLVVLAVLALITLIGCCAYRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGGLVVLAVLALITLIGCCAYRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310

RESULT 15

ID ABG65296
AC ABG65296 standard; Protein; 310 AA.

XX ABG65296;

XX AC

XX ABG65296;

XX 27-AUG-2002 (first entry)

XX Human albumin fusion protein #1971.

DE Albumin fusion protein; therapeutic protein X; human albumin; HA;
KW human serum albumin; HSA; cancer; reproductive disorder;
KW digestive disorder; immune disorder; endocrine disorder;
KW haematopoietic disorder; neural disorder; connective disorder;
KW cytostatic; antiinfectility; antiinflammatory; antiulcer;

KW immunomodulator; anti-HIV; antidiabetic; haemostatic; nootropic;
KW neuroprotective; antiparkinsonian; antimicrobial; neuroleptic;
KW osteopathic; antiarthritic.

XX Homo sapiens.
OS Synthetic.

XX WO200177137-A1.

XX 18-OCT-2001.

XX 12-APR-2001; 2001WO-US11988.

XX 12-APR-2000; 2000US-229358P.

XX 25-APR-2000; 2000US-199384P.

XX 21-DEC-2000; 2000US-256931P.

XX (HUMA-) HUMAN GENOME SCI INC.

XX Rosen CA, Haseltine WA;

XX WPI; 2002-010886/01.

XX New fusion protein for treating disease e.g. diabetes comprises an
PT albumin fused to a therapeutic protein -
XX
XX Claim 1; Page 1893-1894; 2102pp; English.

XX The present invention relates to albumin fusion proteins comprising a
CC therapeutic protein X and human albumin (HA, also known as human serum
CC albumin, HSA). The proteins are useful for treating a disease or
CC disorder that may be modulated by therapeutic protein X. The albumin
CC extends the shelf-life of protein X, and may increase its biological
CC in vitro/in vivo activity. The protein is useful for treating and
CC diagnosing disorders such as cancer, reproductive disorders, digestive
CC disorders (e.g. Crohn's disease, ulcerative colitis), immune disorders
CC (e.g. acquired immunodeficiency syndrome, AIDS), endocrine disorders
CC (e.g. diabetes), haematopoietic disorders, neural disorders
CC (e.g. Alzheimer's, Parkinson's, Creutzfeldt-Jacob disease,
CC encephalomyelitis, meningitis, schizophrenia), and connective disorders
CC (e.g. osteoporosis, arthritis). ABG63326-ABG65518 represent albumin
CC fusion proteins of the invention.

SQ Sequence 310 AA;

Query Match 99.6%; Score 1629; DB 23; Length 310;
Best Local Similarity 99.7%; Pred. No. 7.2e-133;
Matches 309; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MALRRPRLRLCARLPDFFLLLRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
DB 1 MALRRPRLRLCARLPDFFLLLRGCLIGAVNLKSSNRTPVQEFESVELSCIITDSQT 60
QY 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKINWVTRRDSALYRCEVVAR 120
DB 61 SDPRIWKKIODEQTYVFFDNKIQGLAGRAEILGKTSLKINWVTRRDSALYRCEVVAR 120
QY 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
DB 121 NDRKEIDEIVELTVQVKPVPVCRVPKAVPVGKMATLHCQSEGHPRPHYSWYRNDVPL 180
QY 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYCIASNDAGSARCEQMEVYDL 240
DB 181 PTDSRANPRFRNSSHLNSETGLVFTAVHKDDSGQYCIASNDAGSARCEQMEVYDL 240
QY 241 NIGGIIGGLVVLAVLALITLIGCCAYRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300
DB 241 NIGGIIGGLVVLAVLALITLIGCCAYRRGYFINNKQDGESYKPKGPDGVNYIRTDDEG 300
QY 301 DFRHKSSFVI 310
DB 301 DFRHKSSFVI 310

Search completed: December 15, 2003, 14:51:09
Job time : 43 secs
